



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Alcavis International, Inc.  
% Mr. Gary J. Mishkin, M.S.  
VP, Research and Development  
8322 Helgerman Court  
Gaithersburg, Maryland 20877

FEB 15 2007

Re: K061467

Trade/Device Name: ExSept WC Wound Cleanser  
Regulatory Class: Unclassified  
Product Code: FRO  
Dated: December 22, 2006  
Received: December 26, 2006

Dear Mr. Mishkin:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

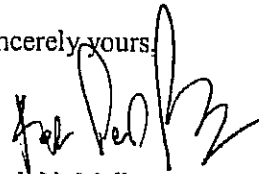
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

Page 2 – Mr. Gary J. Mishkin, M.S.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0115. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Mark N. Melkerson  
Director  
Division of General, Restorative  
and Neurological Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

1061467

## Indications for Use

510(k) Number (if known):

Device Name: ExSept WC Wound Cleanser

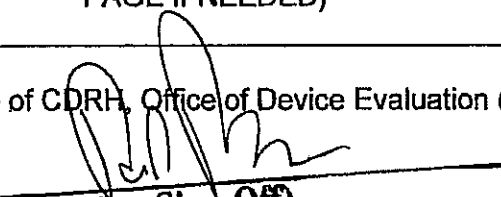
### Indications For Use:

For debridement and the removal of foreign material and debris from exudating and/or dirty wounds, abrasions and minor irritations, cuts, exit sites, and intact skin.

Prescription Use   X   AND/OR Over-The-Counter Use         
(Part 21 CFR 801 Subpart D) (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER  
PAGE IF NEEDED)

Concurrence of CDHR, Office of Device Evaluation (ODE)

  
(Division Sign-Off)

Division of General, Restorative,  
and Neurological Devices

510(k) Number

1061467

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Device Classification Name	<a href="#">Dressing, Wound, Drug</a>
510(K) Number	K061467
Device Name	EXSEPT WC WOUND CLEANSER
Applicant	ALCAVIS INTERNATIONAL, INC. 8322 Helgerman Court Gaithersburg, MD 20877
Contact	Gary J Mishkin
Classification Product Code	<a href="#">FRO</a>
Date Received	05/26/2006
Decision Date	02/15/2007
Decision	Substantially Equivalent (SE)
Classification Advisory Committee	General & Plastic Surgery
Review Advisory Committee	General & Plastic Surgery
Statement/Summary/Purged Status	Summary Only
Summary	<a href="#">Summary</a>
Type	Traditional
Reviewed By Third Party	No
Expedited Review	No